

DETECTOR AGENT FOR HALOGEN COMPOUNDS AND
METHOD FOR DETECTING HALOGEN COMPOUNDS

[Detailed Description of The Invention]

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[Field of the Invention]

The present invention relates to a detector agent for halogen compounds, which contains curcumin or Bromocresol Green as a discoloring component, and a method for detecting halogen compounds by use of said detector agent, and the present invention is usual to judge the leakage of halogen compounds which are used in a semiconductor-manufacturing plant or the like or the break-through of a solid treatment agent for removal of harmful substances.

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[Prior Arts]

Gases containing halogen compounds such as F₂, Cl₂, ClF₃, HF, BrF₅ and SiCl₄ are useful in many applications, for instance, in a field of chemical industries including a semiconductor-manufacturing plant so as to utilize a higher reactivity of halogen atoms. In a case where they are used, various halogen compounds are produced as their reaction products. These halogen compounds possess in many cases a higher reactivity, in which their danger and toxicity are intensive. Accordingly, it is not permitted to release gases containing these halogen compounds as they are, and it is desired to release them to the atmosphere after they are made harmless by a harmful substances removal apparatus.

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As such a harmful substances removal apparatus as mentioned above, there have been known apparatuses for removing or decomposing halogen compounds through a wet treatment such as an alkali scrubber or a dry treatment using a solid treatment agent such as a decomposition catalyst, solid reaction agent or adsorbent. In a case where a solid treatment agent is used, it is necessary to detect halogen compounds on the downstream side

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thereof so that the deterioration of the solid treatment agent is judged because it has a life time (a break-through time) determined by its catalyst poisoning, reactive deterioration or adsorptive saturation.

As for a method for detecting halogen compounds, a method of using a detector agent which is discolored due to the presence of halogen compounds is simple. There has been known, for example, a detector agent containing a molybdate as a discoloring component for dichlorosilane.

[Problems Sought for Solution by the Invention]

For various halogen compounds which are used in a semiconductor-manufacturing plant or the like, however, there has not been known till now a detector agent which is effectively discolored at a lower concentration. In addition, curcumin or Bromocresol Green has been known only as a discoloring component for the detection of boron or the indication of pH.

It is therefore an object of the present invention to provide a detector agent which is effectively discolored at a lower concentration for various halogen compounds and a method for detecting halogen compounds by use of said detector agent.

[Means for Solution of the Problems]

In order to achieve the aforementioned purpose, the inventors have studied hard the usability of various kinds of coloring compounds, have found a fact that the aforementioned purpose can be achieved by using curcumin or Bromocresol Green as a discoloring component, and have completed the present invention.

Namely, the detector agent according to the present invention is a detector agent for halogen compounds, which contains curcumin as a discoloring component, or a detector agent for halogen compounds, which contains Bromocresol Green as a discoloring component.

In the above-mentioned detector agent, said discoloring component is preferably supported on granular activated alumina.

On the other hand, the detecting method according to the present invention is a method for detecting halogen compounds, which comprises bringing anyone of said detector agents into contact with a gas to be detected that may contain halogen compounds, as said detector agent is disposed on a position where it can be visually observed from the outside.

Furthermore, the detecting method according to the present invention is a method for detecting halogen compounds, which comprises visually observing from the outside anyone of said detector agents disposed on the upper layer of a solid treatment agent for removing or decomposing halogen compounds, as a gas to be treated is caused for treatment to flow from the bottom side of a container filled with said solid treatment agent, whereby halogen compounds are detected to judge the deterioration of said solid treatment agent.

[Operational Effects]

Since curcumin or Bromocresol Green is contained as a discoloring component, according to the detector agent of the present invention, there can be provided a detector agent which is effectively discolored at a lower concentration for various halogen compounds, as the results of Examples exhibit.

In a case where said discoloring component is supported on granular activated alumina, the detecting sensitivity thereof is enhanced because the effective contact area of the discoloring component is increased and it becomes easy to visually observe the change in color tone. Furthermore, the fluidity of a gas to be detected becomes well.

Since such a detector agent as mentioned above is brought into contact with a gas to be detected that may contain halogen compounds, as said detector agent is disposed on a position where it can be visually observed from the outside, according to a detecting method of the present invention, on the other hand, an effective detection at a lower concentration can be performed for various halogen compounds.

Since various halogen compounds contained in a gas left after treatment can be effectively detected at a lower concentration by such a simple method that a detector agent is disposed on the upper layer of a solid treatment agent, as mentioned above, according to another detecting method of the present invention, the deterioration of said solid treatment agent can be securely judged.

[Embodiments of the Invention]

Embodiments of the present invention will be described below in an order of the detector agent for halogen compounds and the method for detecting halogen compounds.

(Detector agent for halogen compounds)

As halogen compounds which can be detected in the present invention, there are exemplified SiH_2Cl_2 , HF, F_2 , HBr, Cl_2 , ClF_3 , TiCl_4 , BCl_3 , HI and etc., in a case where curcumin is contained as a discoloring component. And further there are exemplified SiH_2Cl_2 , HF, Cl_2 , BCl_3 , SiHCl_3 , BF_3 , SiF_4 , SiCl_4 , WF_6 and etc., in a case where Bromocresol Green is contained as a discoloring component.

Curcumin which is one kind of diketones has a chemical formula of $\text{C}_{21}\text{H}_{20}\text{O}_6$ and is orange crystal having a melting point of 183°C , and it is used as a discoloring component for the detection of boron or the indication of pH. Curcumin exhibits a yellowish orange appearance when it is supported in a proper amount on a carrier.

Bromocresol Green has a chemical formula of $\text{C}_{21}\text{H}_{14}\text{Br}_4\text{O}_5\text{S}$ and is colorless crystal having a melting point of 217°C , and it is used as a discoloring component for the indication of pH. Bromocresol Green exhibits a blue appearance when it is supported in a proper amount on a carrier such as activated alumine.

It is preferable, for using both these discoloring components as a detector agent, to support or deposit them on a granular material or the like,

and more preferable to use them as they are supported on a granular porous carrier. Although various materials such as alumina, silica gel and diatomaceous earth can be used as the carrier, white materials are preferable and granular activated alumina is preferred because of the above-mentioned reasons. In addition, the grain size of such granular carriers is preferably 2 ~ 5mm.

As for a method of supporting said discoloring component on a carrier, there are used any methods. Namely, such discoloring component can be supported on a carrier, for instance, by a conventional known method which comprises dissolving curcumin or Bromocresol Green in a suitable solvent such as alcohol, immersing a carrier in the resulting mixture, and then drying the carrier to remove the solvent. In a case where a discoloring component is used as it is supported on a carrier, the supporting quantity is preferably 0.001 ~ 0.1 wt%. If the supporting quantity becomes too larger, the discoloring becomes difficult to judge, and if the supporting quantity becomes contrarily too smaller, the discoloring is not almost seen even when the discoloring component is contacted with halogen compounds.

Furthermore, a discoloring component may be supported or deposited on fibers, a fiber cloth or a porous membrane. In this case, the supporting or deposition of a discoloring component will be satisfactorily carried out by impregnating or coating a solution of a discoloring component in or on these carriers and then drying them to remove a solvent, similarly to the case of the above-mentioned carriers. In order to be prevented from being discolored, in addition, the detector agent of the present invention is preferably preserved as it is protected from moisture and direct sunlight, for instance, by being put in a vinyl bag.

(Method for detecting halogen compounds)

The detecting method according to the present invention comprises bringing a detector agent of the present invention into contact with a gas to be detected that may contain halogen compounds, as the detector agent is

disposed on a position where it can be visually observed from the outside.
Concretely, this detecting method will be satisfactorily carried out by charging
said detector agent in a drum (column) which is transparent at least in part, as
in a case of such kind of detector agent of the prior art, and causing a gas to
5 be detected, for example a discharge gas which has been subjected to a
harmful substances removal treatment, to flow at a proper speed through said
column.

While the temperature condition on contact may be normal temperature
at that time, the detector agent is usable also at an elevated temperature (for
10 example, 60 ~ 80°C) in the harmful substances treatment. Since the
discoloring is caused occasionally when moisture is contained in a gas to be
detected, in addition, the gas to be detected may be previously treated by a
drying agent or the like. Further, the column is preferably disposed by
excepting a position where sunlight is directly irradiated thereto.

15 A preferred embodiment of to the present invention comprises, as
shown in Fig.1, visually observing from the outside a detector agent 3
disposed on the upper layer of a solid treatment agent 1 for removing or
decomposing halogen compounds, as a gas to be treated is caused for
treatment to flow from the bottom side of a container 2 filled with said solid
20 treatment agent 1, whereby halogen compounds are detected to judge the
deterioration of said solid treatment agent 1.

In the inside of said container 2 is provided a support plate 4
possessing a dispersing function for the gas to be treated and a supporting
function for the solid treatment agent 1, and on the peripheral wall of the
25 container 2 is provided an inspection hole 2a, through which the detector
agent 3 can be visually observed from the outside. And further, a heating
means (not shown) for heating the solid treatment agent 1 to a temperature
suitable for the removal or decomposition of halogen compounds is provided
round the periphery of the container 2 or in the inside thereof, as necessary.

30 As the solid treatment agent 1, there can be used anyone of known
decomposition catalysts, solid reaction agents, adsorbents and the likes which

remove or decompose halogen compounds. The solid treatment agent 1 is properly heated to a temperature depending on its kind by the heating means, and the quantity (the charging height) of the solid treatment agent 1 and the feed flow rate of a gas to be treated are properly set.

5 The charging height of the detector agent 3 is preferably more than 50mm, and the discharge quantity of halogen compounds can be estimated in a some extent in accordance with the discoloring speed of the detector agent 3. Accordingly, the solid treatment agent 1 can be exchanged at a suitable point of time, by previously making the exchanging point of time due to the
10 deterioration of the solid treatment agent 1 to correspond to the discoloring speed of the detector agent 3 depending on the kind or the like of the solid treatment agent 1. Moreover, the concentration of halogen compounds in a discharge gas can be maintained at a concentration lower than TLV value, by predetermining the height of the discoloring as a reference of the odor
15 allowable concentration (TLV value) based on ACGIH in 1988.

[Examples]

In the next place, examples which concretely exhibit the composition and effects of the present invention will be described. .

Example 1:

0.5g of powdery curcumin was mixed and stirred in 2.5L of ethanol so as to be dissolved therein and 5Kg of activated alumina with a diameter of about 4mm were further mixed therein, and the resulting mixture was
25 continuously stirred until the activated alumina was homogenously colored with an orange color. This colored activated alumina was separated from the liquid and dried at about 60°C in a drier so that ethanol remaining therein was sufficiently removed. Thus, a yellowish orange detector agent containing curcumin as a discoloring component (at the supporting quantity of 0.01 wt%)
30 was prepared.

This detector agent was charged to a height of about 50mm in a transparent glass tube (with an inner diameter of 50mm). Gases (at normal temperature) containing halogen compounds given in Table 1 at different concentrations were fed thereto in turn at 1 L/min. The discoloring was visually observed to examine the color left after the discoloring and the minimum concentration, at that the discoloring was caused. And the results are given in Table 1.

[Table 1]

Halogen Compounds	Coloration	Detecting Sensitivity (ppm)
SiH ₂ Cl ₂	Thick green	5
HF	Yellow	3
F ₂	White	1
HBr	White	3
Cl ₂	White	0.5

Example 2 (Bromocresol Green):

1g of powdery Bromocresol Green was mixed and stirred in 2.5L of ethanol so as to be dissolved therein and 5Kg of activated alumina with a diameter of about 4mm were further mixed therein, and the resulting mixture was continuously stirred until the activated alumina was homogenously colored with a blue color. This colored activated alumina was separated from the liquid and dried at normal temperature so that ethanol remaining therein was removed. Thus, a blue detector agent containing Bromocresol Green as a discoloring component (at the supporting quantity of 0.02 wt%) was prepared.

This detector agent was charged to a height of about 50mm in a transparent glass tube (with an inner diameter of 50mm). Gases (at normal temperature) containing halogen compounds given in Table 2 at different concentrations were fed thereto in turn at 1 L/min. The discoloring was visually observed to examine the color left after the discoloring and the minimum

concentration, at that the discoloring was caused. And the results are given in Table 2.

[Table 2]

Halogen Compounds	Coloration	Detecting Sensitivity (ppm)
SiH ₂ Cl ₂	White	5
HF	White	3
Cl ₂	White	0.5
BCl ₃	White	5
SiHCl ₃	White	5
BF ₃	White	1

Example 3:

A solid treatment agent was charged to a height of 300mm in such a harmful substances removal column as shown in Fig.1, which had an inner diameter of 150mm, and the detector agent obtained in Example 1 was charged to a height of about 50mm on the upper layer thereof. A gas to be treated, which contained 2,000 ppm of F₂ in nitrogen gas, was caused to flow at 20 L/min. from the lower portion of the harmful substances removal column, as the detector agent was visually observed from the outside, whereby a treatment of reacting and solidifying F₂ at normal temperature was performed.

After the elapse of about 90 hours, the detector agent began to discolor to white color from the lower portion of its charged layer, and at a point of time when it was discolored up to a position of 80%, the concentration of F₂ in a discharge gas was measured by a F₂ detector based on the diaphragm electrode method. As a result, the concentration of F₂ was 0 ~ 0.5ppm and this was a value lower than TLV value. At this point of time, it could be judged that the capability of the solid treatment agent was almost saturated.

[Brief Description of the Drawings]

Fig.1 is a front view showing one example of the apparatus used in the detecting method according to the present invention.

[Description of Reference Numerals]

5 1 - solid treatment agent, 2 - container, 2a - inspection hole, 3 - detector agent.

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